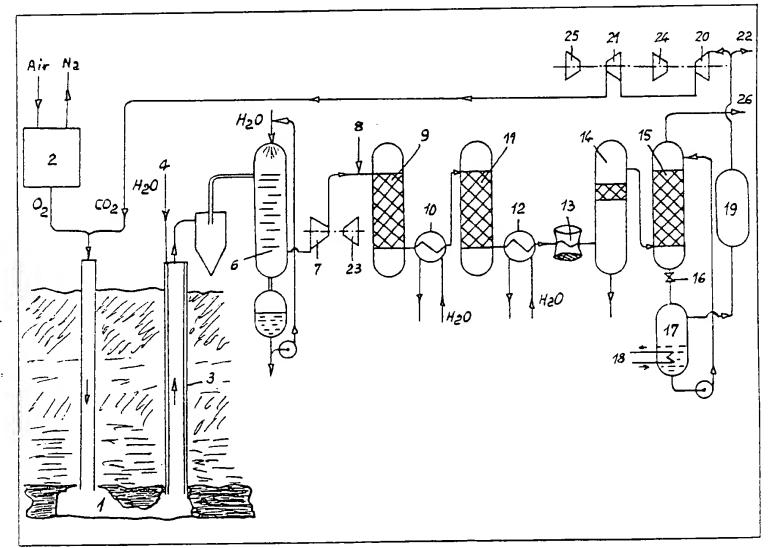
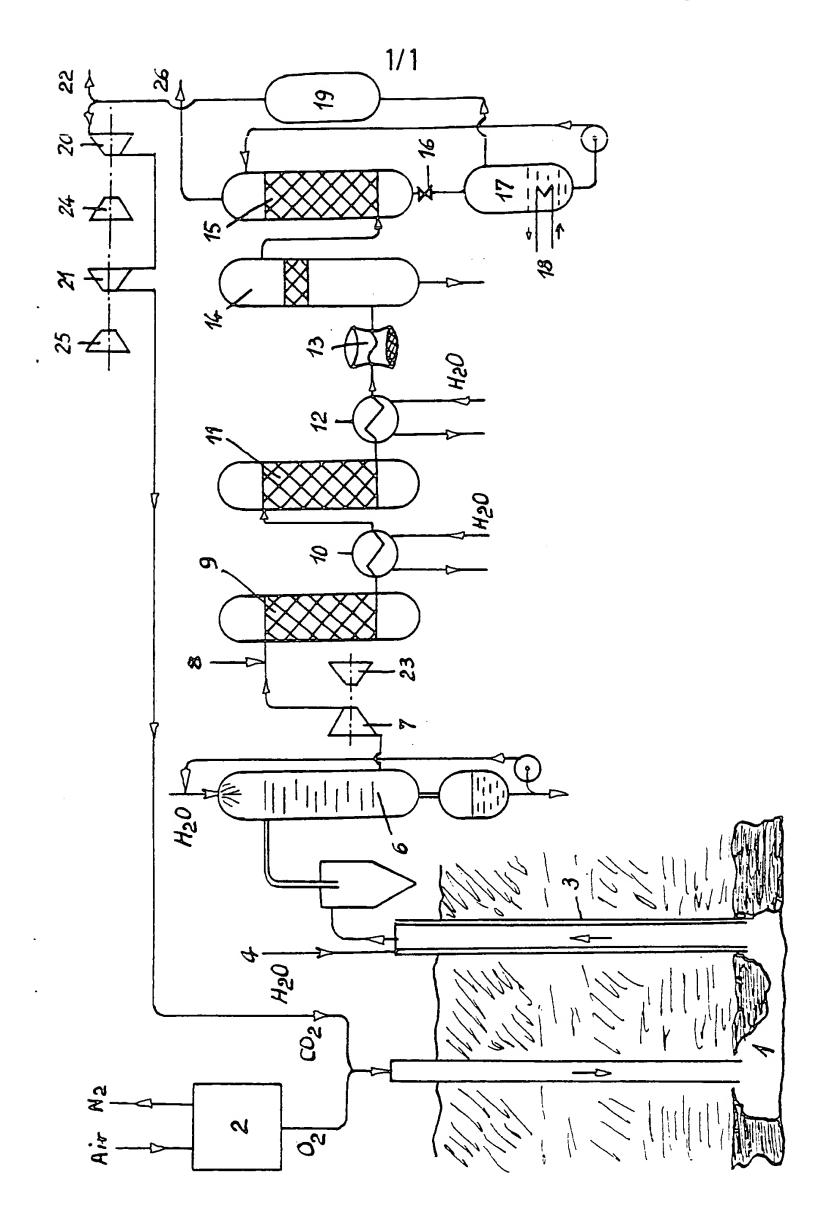
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- (54) Method of producing a gas with a high hydrogen content by subterranean gasification of coal
- (57) A method for the production of a hydrogen rich gas by underground gasification of coal, characterised in that it consists of associating an underground coal gasification operation carried out by means of

oxygen and CO₂, the CO₂ being recovered in the plant in which the gas produced is cleansed, with an operation to cool the crude gas by the injection of water into the bores through which the gas producer discharges and an operation to convert the CO to CO₂, carried out on the surface, employing the water vapour produced during cooling of the gas.



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Method of producing a gas with a high hydrogen content by subterranean gasification of coal

All the extracted coal gasification processes which aim to produce a gas with a high hydrogen content are based on the reaction of the gas with water which can be expressed as: 5 $5 H_2O + C=CO + H_2 - 28.4 \text{ kcal/mole.}$ Generally, the heat required for this reaction to take place is obtained by combustion of a portion of the batch. In older processes, periodic inversions were carried out with the alternate injection of air and steam. More recent processes employ continuous blowing using a gasifying agent consisting of a mixture 10 10 of vapour and oxygen. The same gasifying mixture has been used in the U.S.S.R. and in the U.S.A. during early attempts to produce a gas with a high hydrogen content performed by subterranean gasification of coal at a relatively low pressure in coal deposits located at depths no greater than 300 m. If the production of a hydrogen rich gas by subterranean gasification of mineral coal deposits 15 15 located at a greater depth is envisaged (beyond 700 or 800 m), the costing of the process makes it necessary to increase the gasification pressure to a minimum level of around 30 to 40 bars. Under these conditions of working at great depth and under high pressure, the use of an oxygen vapour mixture as a gasifying agent in which the proportion of vapour may be as much as 65 to 85% is 20 20 not without its drawbacks. To avoid any risk of condensation of the vapour, the temperature of the mixture must at all points remain above a minimum level of around 250°C. This high temperature makes it necessary to reduce the length of the lines used for injecting the gasifying agent and of inserting here and there expansion compensators. Under these conditions, it is 25 virtually out of the question to use a gasification method in which the gasifying agent would be carried 25 through galleries cut by conventional mining techniques and the injection of the gasifying agent by vertical bores providing direct access to the underground gas producer can only be carried out by means of relatively large diameter bores inside of which a heatproof tube is installed. Underground gasification employing an oxygen vapour mixture entails two other types of 30 30 drawback. By reason of its relatively high temperature, the gasifying agent cannot preheat on contact with the rocks which surround the gas producer and it may, on the contrary, give off to them a part of its substantial heat which will proportionately reduce the efficiency of the gas producer in terms of energy. Furthermore, in all underground gasification processes there is an interference between the 35 performance of gasification reactions and the performance of coal pyrolysis reactions, the volatile 35 substances which are released in the depths of the vein having no means of escape other than the gassolid contact surface along which the gasification reactions take place. The release of these volatile substances which contain considerable quantities of hydrogen has a marked inhibiting effect on the reaction of the gas with water; the result is a reduction in the degree of 40 40 decomposition of the vapour injected and a degradation of the gasification efficiency. The object of the process according to the invention is to remedy these various drawbacks. To arrive at this end, the process combines an operation for underground gasification of coal which is carried out by means of a mixture of oxygen and CO2, an operation to cool the crude gas by injection of water into the borings through which the gas producer discharges and a CO conversion 45 operation carried out at the surface employing the steam produced during cooling of the gas. 45 The process according to the invention is likewise characterised by the fact that the CO2 needed to carry out the underground gasification is recovered in the installation in which the gas produced is purified, and in that the reaction heat released during the course of the CO conversion operation is used for producing vapour which is employed in a condensation cycle in order to produce a part of the energy 50 50 needed for operation of the plant. The process according to the invention is described hereinafter with reference to a diagram in the appended drawings. The underground gas producer 1 is supplied with a gasifying agent injected at ambient temperature and at elevated pressure (for example 40 bars to 50 bars). This gasifying agent consists of 55 a mixture of oxygen originating from the plant 2 for separating oxygen from the air and CO₂ emanating 55 from the factory which processes the gas produced. The crude gas arrives at the discharge from the gas producer at a temperature of around 600 to 800°C. It is discharged to the surface via one or more gas bores such as 3, each of which comprises a

expandable towards the bottom.

A line 4 provided with suitable regulating devices injects water under pressure into the annular space which separates the lining from the inner tube. This water evaporates on contact with the wall of the inner tube and the resultant vapour blends with the crude gas at the foot of the shaft.

60 metal lining cemented into the soil and an inner metal tube suspended from the shaft head and freely

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decomposition of the volatile matter in the coal.

Applying this model to a deposit of anthracite-bearing coal containing 7% volatile matter in relation to pure coal, the following results are found:

55 CO₂ + C reactions and on the hypothesis that the methane produced emanates substantially from

A calculation model is employed which is based on the conventional balances of $H_2O + C$ and

I. Gasification by an oxygen-vapour mixture

Composition of the gas (% by volume of crude gas)

	Gas	Crude at 900°C	Cooled to 200°C	Converted	Cleansed	
5	CO :	33.5	33.5	23.0	23.0	5
	CO ₂ :	15.5	15.5	26.0		
	H ₂ :	36.0	36.0	46.5	46.5	
	CH ₄ :	2.5	2.5	2.5	2.5	
	H₂O :	12.5	62.5	52.0		
	•				70.0	10
10		100.0	150.0	150.0	72.0	10

Under these working conditions, the gasification efficiency (P.C.I. of the crude gas/P.C.I. of the gasified coal) is as much as 88%.

The consumption of gasifying agents amounts to:

— 0.175 mole of oxygen and

— 0.407 mole of vapour per mole of crude gas produced. 15

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II. Gasification employing an oxygen-CO₂ mixture

Composition of the gas (in % by volume of crude gas)

	Gas	Crude at 900°C	Cooled at 200°C	Converted	Cleansed	
20	CO :	55.0	55.0	20.0	20.0	20
	CO ₂ :	37.0	37.0	72.0		
	- H ₂ :	5.5	5.5	40.5	40.5	
•	- CH₄ :	2.2	2.2	2.2	2.2	
	H ₂ O :	0.3	50.3	15.3		
25	•	100.0	150.0	150.0	62.7	25

The gasification efficiency (P.C.I. of the crude gas/P.C.I. of the gasified coal) is 86%.

The consumption of gasifying agents amounts to:

- 0.187 mole of oxygen and

— 0.441 mole of CO₂ per mole of crude gas produced.

if these figures for gasification efficiency and consumption of gasifying agents are reduced to one 30 and the same production of cleaned gas, the figures obtained are to the advantage of gasification employing the oxygen-vapour mixture.

Indeed, taking into account the reduction in calorific output resulting from the CO conversion operation, we have:

35 For gasification using oxygen-vapour:

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Gasification efficiency (cleaned gas/coal) = 86.1%

Oxygen consumption:
$$0.175 \times \frac{100}{72} = 0.243$$
 mole/mole

Consumption of vapour: 0.407
$$\times \frac{100}{72} = 0.565$$
 mole/mole

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3**5**

Gasification efficiency (cleaned gas/coal) = 79.3%

$$0xygen consumption: 0.187 \times \frac{100}{62.7} = 0.298 \text{ mole/mole}$$

$$CO_2$$
 consumption: 0.441 $\times \frac{100}{62.7} = 0.703$ mole/mole

5 However, the conclusion is amended if one examines the overall energy balance of the process; indeed, two major factors are to the advantage of the oxygen-CO₂ process:

1) the compression of one mole of CO₂ at 1 bar to 45 bars consumes substantially less energy than the production of one mole of vapour at the same pressure;

2) the heat liberated by the operation to convert the CO is not heat lost, but heat which can be 10 recovered in the form of vapour at a relatively high thermal level (of around 300 to 400°C).

Bearing in mind these two factors and if one attributes an efficiency of 40% to the operation of conversion of heat energy into mechanical energy, the final energy balance looks like this:

Gasification using oxygen-vapour

15		kcal/cu.m. N clean gas	% of P.C.I. of gasified coal	15
	Energy content of the gas:	2,927	86.1	-
	Heat recovery from conversion:	+ 64	+ 1.9	
	Production of oxygen and vapour:	-630	-18.5	
	Compression of the gas prior to conversion:	-200	- 5.9	
20	Energy efficiency	2,161	63.6	20

Gasification using oxygen-CO2

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	kcal/cu.m. N clean gas	% of P.C.I. of gasified coal	
Energy content of the gas:	2,927	79.3	_
Heat recovery from conversion:	+246	+ 6.6	25
Production of oxygen and compression of CO ₂ :	-610	-16.5	
Compression of the gas prior to conversion:	-230	- 6.2	
Energy efficiency	2,333	63.2	

These results show that from the point of view of energy efficiency the two processes are virtually 30 the same and this conclusion would be further reinforced if separation of the CO₂ in the separator 17 were to take place at optimum pressure, in excess of atmospheric pressure, and if one were to take into account the inhibiting effect of the liberation of volatile matter from the coal on the gas reaction to water.

To sum up, the process according to the invention can benefit from all the advantages made 35 available by employing a non-condensable gasifying agent which can be used at ambient temperature without involving any disadvantages with regard to the energy efficiency of the plant. This result is obtained by an association of the operations of underground gasification of the coal, cooling of the gas and conversion of the CO which makes it possible chemically to exploit the inevitable vapour produced by cooling of the crude gas under pressure and temperature conditions which are too low to allow it to 40 be used to advantage in a thermodynamic cycle and which makes it possible to exploit the heat released 40 by the conversion reaction, producing steam at a relatively high temperature which can be employed in a condensation cycle to produce some of the energy needed for the plant to operate.

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The energy economics of the process are likewise favoured by the fact that the CO_2 which results from cleansing of the gas can be employed as a gasifying agent and that the compression of the CO_2 consumes less energy than the production of an equivalent quantity of water vapour or steam.

CLAIMS

1. A method for the production of a hydrogen rich gas by underground gasification of coal, characterised in that it consists of associating an underground coal gasification operation carried out by means of oxygen and CO₂, the CO₂ being recovered in the plant in which the gas produced is cleansed, an operation to cool the crude gas by the injection of water into the bores through which the gas producer discharges and an operation to convert the CO₂, carried out on the surface, employing the water vapour produced during cooling of the gas.

2. A method for producing a hydrogen rich gas according to Claim 1, characterised in that the reaction heat released during the course of the CO conversion operation is used for producing the vapour which is employed in a condensation cycle in order to produce some of the energy required for operation of the plant.

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